

Entrez PubMed Nucleotide Protein Genome Structure OMIM PMC Journals

Search PubMed for egfr Go Clear

☒ Limits Preview/Index History Clipboard

About Entrez

Text Version

Entrez PubMed

Overview

Help | FAQ

Tutorial

New/Noteworthy

E-Utilities

PubMed Services

Journals Database

MeSH Database

Single Citation Matcher

Batch Citation Matcher

Clinical Queries

LinkOut

Cubby

Related Resources

Order Documents

NLM Gateway

TOXNET

Consumer Health

Clinical Alerts

ClinicalTrials.gov

PubMed Central

Display Summary Show: 100 Sort Send to Te

Items 1-12 of 12

- ☐ 1: [Cohen EE, Rosen F, Stadler WM, Recant W, Stenson K, Huo D, Vokes EE.](#) Related



Phase II trial of ZD1839 in recurrent or metastatic squamous cell carcinoma of the head and neck.

J Clin Oncol. 2003 May 15;21(10):1980-7.

PMID: 12743152 [PubMed - indexed for MEDLINE]

- ☐ 2: [Baselga J, Rischin D, Ranson M, Calvert H, Raymond E, Kieback DG, Kaye SB, Gianni L, Harris A, Bjork T, Averbuch SD, Feyereislova A, Swaisland H, Rojo F, Albanell J.](#) Related



Phase I safety, pharmacokinetic, and pharmacodynamic trial of ZD1839, a selective oral epidermal growth factor receptor tyrosine kinase inhibitor, in patients with five selected solid tumor types.

J Clin Oncol. 2002 Nov 1;20(21):4292-302.

PMID: 12409327 [PubMed - indexed for MEDLINE]

- ☐ 3: [Herbst RS, Maddox AM, Rothenberg ML, Small EJ, Rubin EH, Baselga J, Rojo F, Hong WK, Swaisland H, Averbuch SD, Ochs J, LoRusso PM.](#) Related



Selective oral epidermal growth factor receptor tyrosine kinase inhibitor ZD1839 is generally well-tolerated and has activity in non-small-cell lung cancer and other solid tumors: results of a phase I trial.

J Clin Oncol. 2002 Sep 15;20(18):3815-25.

PMID: 12228201 [PubMed - indexed for MEDLINE]

- ☐ 4: [\[No authors listed\]](#) Related



OSI Pharmaceuticals, Genentech and Roche announce data from clinical trial of Tarceva.

Expert Rev Anticancer Ther. 2001 Jun;1(1):4-5. No abstract available.

PMID: 12113131 [PubMed - indexed for MEDLINE]

- ☐ 5: [Dittrich Ch, Greim G, Borner M, Weigang-Kohler K, Huisman H, Amelsberg A, Ehret A, Wanders J, Hanauske A, Fumoleau P.](#) Related




Phase I and pharmacokinetic study of BIBX 1382 BS, an epidermal growth factor receptor (EGFR) inhibitor, given in a continuous daily oral administration.


Eur J Cancer. 2002 May;38(8):1072-80.

PMID: 12008195 [PubMed - indexed for MEDLINE]


- ☐ 6: Ranson M, Hammond LA, Ferry D, Kris M, Tullo A, Murray PL, Miller V, Averbuch S, Ochs J, Morris C, Feyereislova A, Swaisland H, Rowinsky EK. Related

 ZD1839, a selective oral epidermal growth factor receptor-tyrosine kinase inhibitor, is well tolerated and active in patients with solid, malignant results of a phase I trial.
J Clin Oncol. 2002 May 1;20(9):2240-50.
PMID: 11980995 [PubMed - indexed for MEDLINE]


- ☐ 7: Ellis MJ, Coop A, Singh B, Mauriac L, Llombert-Cussac A, Janicke F, Miller WR, Evans DB, Dugan M, Brady C, Quebe-Fehling E, Borgs M. Related

 Letrozole is more effective neoadjuvant endocrine therapy than tamoxifen in ErbB-1- and/or ErbB-2-positive, estrogen receptor-positive primary breast cancer: evidence from a phase III randomized trial.
J Clin Oncol. 2001 Sep 15;19(18):3808-16.
PMID: 11559718 [PubMed - indexed for MEDLINE]


- ☐ 8: Hidalgo M, Siu LL, Nemunaitis J, Rizzo J, Hammond LA, Takimoto C, Eckhardt SG, Tolcher A, Britten CD, Denis L, Ferrante K, Von Hoff DD, Silberman S, Rowinsky EK. Related

 Phase I and pharmacologic study of OSI-774, an epidermal growth factor receptor tyrosine kinase inhibitor, in patients with advanced solid malignancies.
J Clin Oncol. 2001 Jul 1;19(13):3267-79.
PMID: 11432895 [PubMed - indexed for MEDLINE]


- ☐ 9: Ghosh S, Liu XP, Zheng Y, Uckun FM. Related

 Rational design of potent and selective EGFR tyrosine kinase inhibitors as anticancer agents.
Curr Cancer Drug Targets. 2001 Aug;1(2):129-40. Review.
PMID: 12188886 [PubMed - indexed for MEDLINE]


- ☐ 10: Norman P. Related

 OSI-774 OSI Pharmaceuticals.
Curr Opin Investig Drugs. 2001 Feb;2(2):298-304. Review.
PMID: 11816847 [PubMed - indexed for MEDLINE]

- ☐ 11: Bridges AJ. Related

 Chemical inhibitors of protein kinases.
Chem Rev. 2001 Aug;101(8):2541-72. Review. No abstract available.
PMID: 11749388 [PubMed - indexed for MEDLINE]

- ☐ 12: Arteaga CL, Johnson DH. Related

 Tyrosine kinase inhibitors-ZD1839 (Iressa).
Curr Opin Oncol. 2001 Nov;13(6):491-8. Review.
PMID: 11673690 [PubMed - indexed for MEDLINE]

☐ Display ☐ Summary ☐ Show: ☐ Sort ☐ Send to
Items 1-12 of 12

10/622,787

Thomas McKenzie

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID:sssptal611txm

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

* * * * * Welcome to STN International * * * * *

NEWS 1 Web Page URLs for STN Seminar Schedule - N. America
NEWS 2 "Ask CAS" for self-help around the clock
NEWS 3 May 10 PROUSDDR now available on STN
NEWS 4 May 19 PROUSDDR: One FREE connect hour, per account, in both May
and June 2004
NEWS 5 May 12 EXTEND option available in structure searching
NEWS 6 May 12 Polymer links for the POLYLINK command completed in REGISTRY
NEWS 7 May 17 FRFULL now available on STN
NEWS 8 May 27 New UPM (Update Code Maximum) field for more efficient patent
SDIs in CPlus
NEWS 9 May 27 CPlus super roles and document types searchable in REGISTRY
NEWS 10 May 27 Explore APOLLIT with free connect time in June 2004
NEWS 11 Jun 22 STN Patent Forums to be held July 19-22, 2004
NEWS 12 Jun 28 Additional enzyme-catalyzed reactions added to CASREACT
NEWS 13 Jun 28 ANTE, AQUALINE, BIOENG, CIVILENG, ENVIROENG, MECHENG,
and WATER from CSA now available on STN(R)

NEWS EXPRESS MARCH 31 CURRENT WINDOWS VERSION IS V7.00A, CURRENT
MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
AND CURRENT DISCOVER FILE IS DATED 26 APRIL 2004
NEWS HOURS STN Operating Hours Plus Help Desk Availability
NEWS INTER General Internet Information
NEWS LOGIN Welcome Banner and News Items
NEWS PHONE Direct Dial and Telecommunication Network Access to STN
NEWS WWW CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that
specific topic.

All use of STN is subject to the provisions of the STN Customer
agreement. Please note that this agreement limits use to scientific
research. Use for software development or design or implementation
of commercial gateways or other similar uses is prohibited and may
result in loss of user privileges and other penalties.

* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 13:50:10 ON 06 JUL 2004

=> file reg

FILE 'REGISTRY' ENTERED AT 13:50:24 ON 06 JUL 2004

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.

PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

COPYRIGHT (C) 2004 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 5 JUL 2004 HIGHEST RN 704870-92-8
 DICTIONARY FILE UPDATES: 5 JUL 2004 HIGHEST RN 704870-92-8

TSCA INFORMATION NOW CURRENT THROUGH JANUARY 6, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

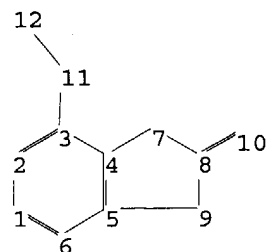
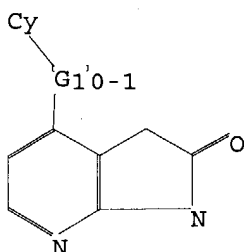
Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:
<http://www.cas.org/ONLINE/DBSS/registryss.html>

=>

Uploading C:\Program Files\Stnexp\Queries\10622787.str

1—2 13—1.2



chain nodes :

10 11 12 13 14

ring nodes :

1 2 3 4 5 6 7 8 9

chain bonds :

3-11 8-10 11-12 13-14

ring bonds :

1-2 1-6 2-3 3-4 4-5 4-7 5-6 5-9 7-8 8-9

exact/norm bonds :

3-11 4-7 5-9 7-8 8-9 8-10 11-12

10/622,787

Thomas McKenzie

exact bonds :

13-14

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6

G1:O,S,N, [*1-*2]

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
11:CLASS 12:Atom 13:CLASS 14:CLASS

L1 STRUCTURE UPLOADED

=> s l1

SAMPLE SEARCH INITIATED 13:50:44 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 476 TO ITERATE

100.0% PROCESSED 476 ITERATIONS
SEARCH TIME: 00.00.01

0 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 8212 TO 10828
PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s l1 full

FULL SEARCH INITIATED 13:50:53 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 9222 TO ITERATE

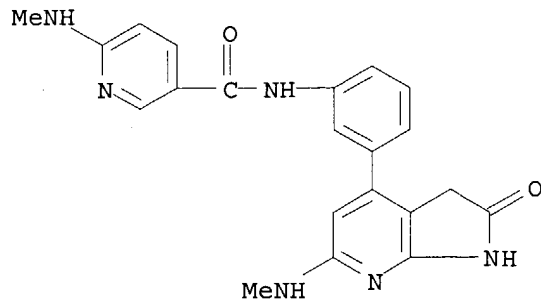
100.0% PROCESSED 9222 ITERATIONS
SEARCH TIME: 00.00.01

3 ANSWERS

L3 3 SEA SSS FUL L1

=> d 1-3 ide cbib pi

L3 ANSWER 1 OF 3 REGISTRY COPYRIGHT 2004 ACS on STN
RN 346599-84-6 REGISTRY
CN 3-Pyridinecarboxamide, N-[3-[2,3-dihydro-6-(methylamino)-2-oxo-1H-pyrrolo[2,3-b]pyridin-4-yl]phenyl]-6-(methylamino)- (9CI) (CA INDEX NAME)
FS 3D CONCORD
MF C21 H20 N6 O2
SR CA
LC STN Files: CA, CAPLUS, USPAT2, USPATFULL
DT.CA Caplus document type: Patent
RL.P Roles from patents: BIOL (Biological study); PREP (Preparation); USES (Uses)



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 135:76886 Preparation of 4-substituted 7-azaindolin-2-ones and their use as protein kinase inhibitors. Liang, Congxin; Sun, Li; Wei, Chung Chen; Tang, Peng Cho; McMahon, Gerald; Hirth, Klaus Peter; Cui, Jingrong (Sugen, Inc., USA). PCT Int. Appl. WO 2001046196 A1 20010628, 97 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2000-US34259 20001221. PRIORITY: US 1999-PV171288 19991221.

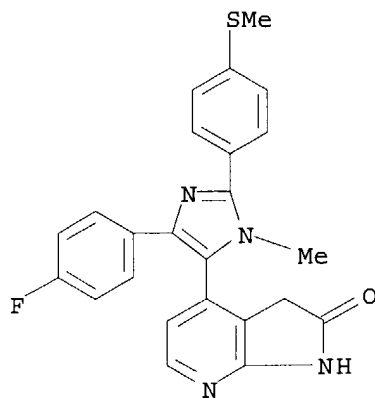
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2001046196	A1	20010628	WO 2000-US34259	20001221
W:		AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM		
RW:		GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		
EP 1244672	A1	20021002	EP 2000-990229	20001221
R:		AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR		
US 2002183319	A1	20021205	US 2000-741737	20001221
US 6610688	B2	20030826		
JP 2003518123	T2	20030603	JP 2001-547106	20001221
US 2004053924	A1	20040318	US 2003-622787	20030721

L3 ANSWER 2 OF 3 REGISTRY COPYRIGHT 2004 ACS on STN
RN 346599-82-4 REGISTRY
CN 2H-Pyrrolo[2,3-b]pyridin-2-one, 4-[4-(4-fluorophenyl)-1-methyl-2-[4-(methylthio)phenyl]-1H-imidazol-5-yl]-1,3-dihydro- (9CI) (CA INDEX NAME)
FS 3D CONCORD
MF C24 H19 F N4 O S
SR CA

LC STN Files: CA, CAPLUS, USPAT2, USPATFULL

DT.CA CAPLUS document type: Patent

RL.P Roles from patents: BIOL (Biological study); PREP (Preparation); USES (Uses)



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

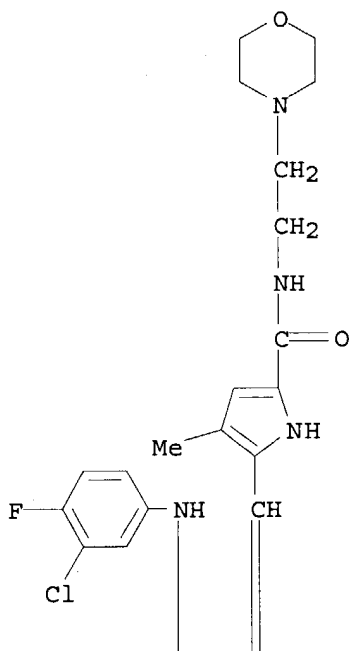
REFERENCE 1: 135:76886 Preparation of 4-substituted 7-azaindolin-2-ones and their use as protein kinase inhibitors. Liang, Congxin; Sun, Li; Wei, Chung Chen; Tang, Peng Cho; McMahon, Gerald; Hirth, Klaus Peter; Cui, Jingrong (Sugen, Inc., USA). PCT Int. Appl. WO 2001046196 A1 20010628, 97 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2000-US34259 20001221. PRIORITY: US 1999-PV171288 19991221.

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001046196	A1	20010628	WO 2000-US34259	20001221

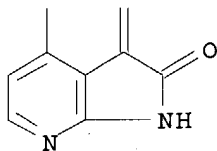
PI	WO 2001046196	A1	20010628	WO 2000-US34259	20001221
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
EP 1244672	A1	20021002	EP 2000-990229	20001221	
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR			
US 2002183319	A1	20021205	US 2000-741737	20001221	
US 6610688	B2	20030826			
JP 2003518123	T2	20030603	JP 2001-547106	20001221	
US 2004053924	A1	20040318	US 2003-622787	20030721	

L3 ANSWER 3 OF 3 REGISTRY COPYRIGHT 2004 ACS on STN
RN 346599-64-2 REGISTRY
CN 1H-Pyrrole-2-carboxamide, 5-[[4-[(3-chloro-4-fluorophenyl)amino]-1,2-dihydro-2-oxo-3H-pyrrolo[2,3-b]pyridin-3-ylidene]methyl]-4-methyl-N-[2-(4-morpholinyl)ethyl]- (9CI) (CA INDEX NAME)
FS 3D CONCORD
MF C26 H26 Cl F N6 O3
SR CA
LC STN Files: CA, CAPLUS, USPAT2, USPATFULL
DT.CA Caplus document type: Patent
RL.P Roles from patents: BIOL (Biological study); PREP (Preparation); USES (Uses)

PAGE 1-A



PAGE 2-A



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 135:76886 Preparation of 4-substituted 7-azaindolin-2-ones and their use as protein kinase inhibitors. Liang, Congxin; Sun, Li; Wei, Chung Chen; Tang, Peng Cho; McMahon, Gerald; Hirth, Klaus Peter; Cui, Jingrong (Sugen, Inc., USA). PCT Int. Appl. WO 2001046196 A1 20010628, 97 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2000-US34259 20001221. PRIORITY: US 1999-PV171288 19991221.

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001046196	A1	20010628	WO 2000-US34259	20001221
EP 1244672	A1	20021002	EP 2000-990229	20001221
US 2002183319	A1	20021205	US 2000-741737	20001221
US 6610688	B2	20030826		
JP 2003518123	T2	20030603	JP 2001-547106	20001221
US 2004053924	A1	20040318	US 2003-622787	20030721

=>

=> logoff

ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF

LOGOFF? (Y)/N/HOLD:.

STN INTERNATIONAL LOGOFF AT 13:52:25 ON 06 JUL 2004

L Number	Hits	Search Text	DB	Time stamp
1	1368	((514/300) or (546/113)).CCLS.	USPAT; US-PGPUB	2004/07/06 14:02
2	28	(((514/300) or (546/113)).CCLS.) and ("pyrrolo[2,3-b]pyridin" OR "pyrrolo[2,3-b]" OR "[2,3-b]pyridin" OR "pyrrolo(2,3-b)pyridin" OR "pyrrolo(2,3-b)" OR "(2,3-b)pyridin" OR "[2,3-b]" OR "(2,3-b)" OR "2,3-b")	USPAT; US-PGPUB	2004/07/06 14:06